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(54) Infrared radiation absorbing glass

Infrarotstrahlung absorbierendes Glas

Verre absorbant rayons infrarouges

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(56) References cited:

EP-A- 0 561 337	EP-A- 0 565 882
EP-A- 0 604 314	EP-A- 0 677 492
EP-A- 0 709 344	EP-A- 0 748 776
US-A- 3 723 142	US-A- 4 701 425
US-A- 5 344 798	US-A- 5 411 922

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Description

FIELD OF THE INVENTION

5 [0001] The present invention relates to an infrared radiation absorbing glass having a high visible light transmission.

BACKGROUND OF THE INVENTION

10 [0002] From the standpoint in reduction of the load of air conditioning in automobiles, a glass having infrared absorbing power has recently been proposed as window panes of automobiles.

[0003] For example, greenish blue-tinted glass having a relatively high Fe_2O_3 content to exhibit improved infrared absorbing power has been developed for use in automobiles. A typical example of such a greenish blue-tinted infrared absorbing glass has a composition comprising, in % by weight: 71% SiO_2 , 1.5% Al_2O_3 , 4% MgO , 8.6% CaO , 13.5% Na_2O , 0.7% K_2O , 0.55% total iron oxides in terms of Fe_2O_3 , and 0.17 FeO in terms of Fe_2O_3 . The glass having this composition with a thickness of 4 mm has a visible light transmission of 78% as measured with the CIE standard illuminant A and a solar radiation transmission of 54%.

15 [0004] Glass having a bronze tint, on the other hand, contains Fe_2O_3 in an amount smaller than that in the greenish blue-tinted glass so as to have fair infrared absorbing power while maintaining visible light transmission. A typical example of infrared radiation absorbing glass having a bronze tint has a composition of, in % by weight: 72% of SiO_2 , 1.5% Al_2O_3 , 4% MgO , 8% CaO , 13.5% Na_2O , 0.7% K_2O , 0.24% total iron oxides in terms of Fe_2O_3 , 0.054% of FeO in terms of Fe_2O_3 , 0.001% CoO , and 0.001 wt% of Se . The glass having this composition with a thickness of 4 mm has a visible light transmission of 78% as measured with the CIE standard illuminant A and a solar radiation transmission of 70%.

20 [0005] In the above-described conventional infrared radiation absorbing glass, its infrared absorbing power is imparted by Fe^{2+} (FeO). However, if the FeO concentration in a greenish blue-tinted glass is increased in order to obtain higher infrared absorbing power, the absorption of FeO in the visible region increases, and as a result, there is particularly a problem that the visible light transmission required of windowpanes of vehicles cannot be obtained. In a glass which forms a color utilizing absorption of Se in the visible region, such as bronze type glass, there is the problem that greater restriction is imposed on the FeO concentration, and a high infrared absorbing power is not obtained.

30 [0006] EP-A-0748776, constituting prior art in accordance with Article 54(3) EPC, discloses an ultraviolet and infrared radiation absorbing glass comprising, as basic glass components,

65 to 80% by weight of SiO_2 ,
 0 to 5% by weight of Al_2O_3 ,
 35 0 to 10% by weight of MgO ,
 5 to 15% by weight of CaO ,
 10 to 18% by weight of Na_2O ,
 0 to 5% by weight of K_2O ,
 5 to 15% by weight in total of $\text{MgO} + \text{CaO}$,
 40 10 to 20% by weight in total of $\text{Na}_2\text{O} + \text{K}_2\text{O}$, and
 0 to 5% by weight of B_2O_3 ; and

as coloring components,

45 0.75 to 0.95% by weight, in terms of Fe_2O_3 , of total iron oxide,.
 1.2 to 1.4% by weight of CeO_2 , and
 0 to 0.5% by weight of TiO_2 ,

and satisfying the following equation:

$$50 \quad (0.2 \times (\text{CeO}_2) - 0.04) \leq (\text{FeO/T-Fe}_2\text{O}_3) \leq (0.2 \times (\text{CeO}_2) + 0.08)$$

55 wherein (CeO_2) represents the amount of CeO_2 in terms of percent by weight, and $(\text{FeO/T-Fe}_2\text{O}_3)$ represents the weight ratio of FeO in terms of Fe_2O_3 to the total iron oxide in terms of Fe_2O_3 .

[0007] EP-A-0565882 discloses a neutral gray colored infrared and ultraviolet radiation absorbing glass comprising as coloring components, on a weight basis, 0.10 - 0.55% of total iron expressed as Fe_2O_3 , 0.2 - 0.6% of CeO_2 , 15-35 ppm of CoO and 2-18 ppm of Se .

[0008] US-A-3723142 discloses a neutral gray colored heat-absorbing glass consisting essentially of the following base components in percent by weight: 68 to 75% SiO₂, 0 to 5% Al₂O₃, 5 to 15% CaO, 0 to 10% MgO, 10 to 18% Na₂O, 0 to 5% K₂O, wherein CaO + MgO is 6 to 15% and Na₂O + K₂O is 10 to 20%, together with coloring components of 0.18 to 0.22% Fe₂O₃, 0.0031 to 0.0040% CoO, 0.0006 to 0.0009% Se, and 0-0.0004% NiO.

SUMMARY OF THE INVENTION

[0009] The present invention has been made in view of the above-described problems involved in the prior art.

[0010] An object of the present invention is to provide a method for producing a glass having a high infrared absorbing power and a high visible light transmission.

[0011] According to the present invention, there is provided a method for shifting the absorption peak wavelength in the wavelength range 900-1600 nm of an infrared absorbing glass from less than 1100 nm to 1100 nm or longer, comprising the step of irradiating the glass which comprises by wt.%:

65 to 80% SiO₂,
0 to 5% Al₂O₃,
0 to 10% MgO,
5 to 15% CaO,
10 to 18% Na₂O,
0 to 5% K₂O,
5 to 15% MgO + CaO,
10 to 20% Na₂O + K₂O,
0 to 5% B₂O₃,
0.05 to 1.0% total iron oxide (T-Fe₂O₃) in terms of Fe₂O₃,
0.05 to 2.0% CeO₂,
0 to 1.0% TiO₂,
0 to 0.005% CoO, and
0 to 0.005% Se,

with ultraviolet light to increase the content of FeO in the irradiated glass by reducing Fe(III) to Fe(II), the UV irradiated glass thereby comprising 0.02 to 0.5 wt.% FeO in terms of Fe₂O₃.

[0012] Preferably, the infrared absorbing glass comprises, in % by weight:

0.2 to 0.6% T-Fe₂O₃ in terms of Fe₂O₃,
0.02 to 0.3% FeO in terms of Fe₂O₃,
0.2 to 2.0% CeO₂; and
0.0005 to 0.005% Se,

wherein a relationship between the transmission at 1.050 nm (T₁₀₅₀) and the transmission at 1.200 nm (T₁₂₀₀) is T₁₀₅₀ > T₁₂₀₀.

[0013] Preferably, the infrared absorbing glass comprises, in % by weight, 0.05 to 3% total of one or more components selected from CeO₂, Sb₂O₃, As₂O₃, SnO, and Cu₂O.

[0014] The infrared radiation absorbing glasses of the present invention preferably have a visible light transmission of 70% or more as measured with the CIE standard illuminant A and a total solar energy transmission of less than 70% as measured in the wavelength region of from 300 to 2,100 nm, when the thickness of the glass is 4 mm.

[0015] It is also preferred that the infrared radiation absorbing glasses according to the present invention have a dominant wavelength of 574 to 580 nm as measured with the CIE standard illuminant C and a total sunlight UV transmission of less than 12% as measured in a wavelength region of 297.5 to 377.5 nm according to ISO, when the thickness of the glasses is 4 mm.

[0016] It is further preferred that the infrared radiation absorbing glasses according to the present invention have an ultraviolet transmission of less than 34% at a wavelength of 370 nm, when the thickness of the glasses is 4 mm.

DETAILED DESCRIPTION OF THE INVENTION

[0017] The reasons for limitation of the glass compositions of the infrared radiation absorbing glass according to the present invention are explained below. Hereinafter, unless otherwise indicated, all the percents are by weight.

[0018] Iron oxide is present in glass in the form of Fe₂O₃ (Fe³⁺) and FeO (Fe²⁺). Fe₂O₃ is a component serving to increase an infrared absorbing power, while FeO is a component serving to increase an ultraviolet absorbing power

together with CeO_2 and TiO_2 .

[0019] In general soda lime silica glass composition used as window glass of buildings and vehicles, Fe^{2+} shows its absorption peak at about 1,050 nm and also has an absorption in the visible light region of from 400 to 780 nm. Therefore, if the FeO content is increased, the infrared absorbing power is increased, but a visible light transmission is simultaneously decreased. The gist of the present invention resides in shifting the absorption peak wavelength of Fe^{2+} to a longer wavelength side to minimize the decrease in visible light transmission, thereby achieving both a high infrared absorbing power and a high visible light transmission at a time.

[0020] The absorption peak wavelength of Fe^{2+} is shifted to the longer wavelength side by irradiating the glass with ultraviolet light at around room temperature.

[0021] The inventors of the present invention have found that ultraviolet irradiation of appropriate glass containing Fe_2O_3 under appropriate conditions easily results in formation of Fe^{2+} having its absorption peak wavelength shifted to the longer wavelength side. Specifically, when glass containing a component acting as a photo-reducing agent, such as Ce^{3+} , and Fe^{3+} is irradiated with UV at around room temperature, Fe^{2+} is formed through the reaction: $\text{Fe}^{3+} + \text{e}^- \rightarrow \text{Fe}^{2+}$. While the absorption peak of Fe^{2+} in a general soda lime silica glass composition for windows is at about 1050 nm, that of the Fe^{2+} newly formed in glass on ultraviolet irradiation is shifted to the longer wavelength side, i.e., at about 1,200 nm or even more. This is because the thus formed Fe^{2+} is surrounded by a distorted structure. The absorption peak of glass in an infrared region after ultraviolet irradiation is determined by the overlap between the absorption of the originally present Fe^{2+} before irradiation and the absorption of the newly formed Fe^{2+} after irradiation, and is usually at a wavelength of from 1,100 to 1,250 nm, although varying depending on the glass composition, ultraviolet irradiation conditions, and the like. Ultraviolet irradiation is preferably conducted by using ultraviolet light of 400 nm or less such that an irradiation energy (a product of an irradiation energy density per unit time per unit area and an irradiation time) is 10^5 J/m^2 or more.

[0022] The shift of the absorption peak wavelength to the longer wavelength side can be evaluated by comparing the transmissions at the peak wavelength. In the present invention, the shift is expressed in terms of change in the wavelength at which the transmission has the minimum value in the range of from 900 to 1,600 nm (λ_p) or in terms of the comparison between the transmission at 1,050 nm (T_{1050}) and the transmission at 1,200 nm (T_{1200}). λ_p is obtained, for example, as the minimum value of a transmission curve obtained by subjecting general smoothing treatment to the transmission data measured for each 2 nm or smaller.

[0023] If the total iron oxide ($\text{T-Fe}_2\text{O}_3$) content is too small, the infrared absorption power and ultraviolet absorbing power are decreased. If it is too large, the visible light transmission is decreased. Accordingly, the $\text{T-Fe}_2\text{O}_3$ content is from 0.05 to 1.0%. In the glass composition containing Se, for the formation of a bronze tint, the $\text{T-Fe}_2\text{O}_3$ content is preferably 0.2 to 0.6%.

[0024] If the FeO content is too small, the infrared absorbing power is decreased, and if it is too large, a visible light transmission is decreased. Accordingly, the FeO content is from 0.02 to 0.5%, in terms of Fe_2O_3 . In the glass composition containing Se, for the formation of a bronze tint, the FeO content preferably is from 0.02 to 0.3%.

[0025] CeO_2 is a component for enhancing ultraviolet absorption. It exists in glass in the form of Ce^{3+} or Ce^{4+} . While Ce^{3+} shows weak absorption in the visible region and is more effective for UV absorption, it is an important component for its function as a photo-reducing agent in the present invention. Specifically, when glass containing Ce^{3+} is irradiated with ultraviolet light at around room temperature, Ce^{3+} acts as a photo-reducing agent, emitting an electron through the reaction: $\text{Ce}^{3+} \rightarrow \text{Ce}^{4+} + \text{e}^-$. The thus emitted electron functions in the reaction: $\text{Fe}^{3+} + \text{e}^- \rightarrow \text{Fe}^{2+}$, in which Fe_2O_3 is reduced to form FeO. As described above, the thus formed FeO has its infrared absorption peak at a wavelength of about 1,200 nm or longer, which is a wavelength shifted to the long wavelength side as compared with about 1,050 nm of the FeO formed by a melt method in the production of general soda lime silica glass for windows. As a result, the glass after ultraviolet irradiation has an absorption peak in the infrared region at a wavelength of from about 1,100 to 1,250 nm, and thus can exhibit improved infrared absorbing power while minimizing the decrease in visible light transmission. The CeO_2 content should be 2.0% or less. If the CeO_2 content is too large, the absorption in the short wavelength side in the visible region is so large that the resulting glass assumes a yellow tint. To ensure a satisfactory function as a photo-reducing agent and to obtain an ultraviolet absorbing power, the CeO_2 content is 0.05 to 2.0%, and more preferably 0.2 to 2.0%.

[0026] SiO_2 is a main component for forming a skeleton of the glass. If the SiO_2 content is less than 65%, the glass has a reduced durability. If it exceeds 80%, it is difficult to melt the glass composition.

[0027] Al_2O_3 serves to improve durability of glass. If the Al_2O_3 content exceeds 5%, it is difficult to melt the glass composition. A preferred Al_2O_3 content is from 0.1 to 2%.

[0028] MgO and CaO both serve to improve durability of glass and also to control the devitrification temperature and viscosity in forming the glass. If the MgO content exceeds 10%, the devitrification temperature rises. If the CaO content is less than 5% or higher than 15%, the devitrification temperature rises. If the total content of MgO and CaO is less than 5%, the resulting glass has reduced durability, if the total content exceeds 15%, the devitrification temperature rises.

[0029] Na_2O and K_2O are used as a glass melting accelerator. If the Na_2O content is less than 10%, or if the total content of Na_2O and K_2O is less than 10%, the effect of melting acceleration is weak. If Na_2O content exceeds 18%, or if the total content of Na_2O and K_2O exceeds 20%, durability of glass is decreased. K_2O is effective to enhance the pink color development of Se and simultaneously to increase the ultraviolet absorbing power. It is unfavorable for the K_2O content to exceed 5% because it is more expensive than Na_2O .

[0030] While B_2O_3 is a component used for the improvement of durability of glass and also as a melting aid, it also functions to enhance ultraviolet absorption. If the B_2O_3 content exceeds 5.0%, decrease in the light transmission in the ultraviolet region is also extended to the visible region, and the glass tends to assume a yellow tint. Moreover, disadvantages occur due to vaporization of B_2O_3 in forming glass. Accordingly, the upper limit of the B_2O_3 content should be 5.0%.

[0031] TiO_2 is a component for increasing ultraviolet absorbing power through the mutual action especially with FeO . If the TiO_2 content exceeds 1.0%, the glass assumes a yellow tint.

[0032] CoO is a component which develops a bronze color in the presence of Se. If the CoO content exceeds 0.005%, the visible light transmission is reduced.

[0033] Se develops a pink tint to thereby provide a bronze tint coupled with CoO which has a complementary color to the pink tint. If the Se content is less than 0.0005%, a desired color cannot be obtained. If it exceeds 0.005%, the visible light transmission is reduced.

[0034] Sb_2O_3 , As_2O_3 , SnO , and Cu_2O function as a photo-reducing agent similar to CeO_2 or serve to accelerate the reducing action of CeO_2 . If the total content of any one or more of these 5 components (inclusive of CeO_2) is less than 0.05%, the photo-reducing effect is small. The upper limit of the total content of the 5 components is 3% because of their expensiveness.

[0035] So long as the effect as aimed at in each embodiment of the present invention is not impaired, the glass composition according to the present invention may further contain one or more of ZnO , Bi_2O_3 , NiO , MnO , V_2O_5 , and MoO_3 in a total amount of up to 1%; one or more of F, Cl, and Br in a total amount of up to 1%; and up to 1% of S in terms of SO_3 .

[0036] The present invention is described in greater detail below by reference to the following Examples, but it should be understood that the invention is not construed as being limited thereto.

EXAMPLES 1 TO 5 AND COMPARATIVE EXAMPLES 1 TO 4

[0037] Siliceous sand, dolomite, limestone, soda ash, potassium carbonate, boron oxide, borax, ferric oxide, titanium oxide, cerium oxide, cobalt oxide, sodium selenite, and a carbonaceous material as a reducing agent were compounded at an appropriate ratio so as to obtain a prescribed glass composition. The resulting blend was melted at 1,500°C for 4 hours in an electric furnace, and the molten glass was cast on a stainless steel plate and gradually cooled to room temperature to obtain a glass plate having a thickness of about 6 mm. This glass plate was polished to a thickness of 4 mm to prepare a glass sample.

[0038] A visible light transmission (YA) measured with the CIE standard illuminant A, a total solar energy transmission (TG), an ultraviolet transmission (T_{UV}) specified by ISO, and a dominant wavelength (D_w) and an excitation purity (Pe) measured with the CIE standard illuminant C of the resulting glass sample were obtained. As an additional measure for ultraviolet transmission, an ultraviolet transmission at 370 nm (T_{370}), at which the change in transmission sensitively appears in the course of the steep rise from the absorption end of the transmission curve, was also measured.

[0039] The sample was then irradiated with ultraviolet light of 400 nm or shorter using a sunshine carbon arc lamp at an energy density of $1.43 \times 10^6 \text{ J/m}^2/\text{hr}$ for a prescribed irradiation time to give a prescribed UV energy Q (J/m^2), a product of the energy density and an irradiation time. After the ultraviolet irradiation, the above-described optical characteristics were measured again. The change in absorption peak wavelength of FeO due to ultraviolet irradiation was evaluated from the transmissions at 1,050 nm and 1,200 nm (T_{1050} , T_{1200}) and the wavelength at which the transmission has the minimum value in the range of from 900 to 1,600 nm (λ_p).

[0040] The concentrations of each component and optical characteristics of the ultraviolet irradiated samples and the ultraviolet irradiation energy (Examples 1 to 5) are shown in Table 1 below.

[0041] The concentrations of each component and optical characteristics of the samples before ultraviolet irradiation (Comparative Examples 1 to 4) are shown in Table 2 below. Comparative Example 1 corresponds to Example 1; Comparative Example 2 to Example 2; Comparative Example 3 to Examples 3 and 4; and Comparative Example 4 to Example 5.

[0042] All the concentrations shown in the Tables are expressed in % by weight.

TABLE 1

	Example 1	Example 2	Example 3	Example 4	Example 5
Composition (wt%):					
SiO ₂	71.3	70.5	70.5	70.5	70.5
B ₂ O ₃	0	1.0	0	0	0
Al ₂ O ₃	1.6	1.5	1.5	1.5	1.5
MgO	3.64	4.0	4.0	4.0	3.46
CaO	7.8	8.0	8.0	8.0	8.0
Na ₂ O	12.5	13.0	13.0	13.0	13.0
K ₂ O	0.6	0.7	0.7	0.7	1.3
CeO ₂	1.6	1.55	1.75	1.75	1.7
TiO ₂	0.33	0.03	0.23	0.23	0.10
CoO	0	0	0.0011	0.0011	0.0017
SeO	0	0	0.0011	0.0011	0.0011
T-Fe ₂ O ₃	0.63	0.7	0.35	0.35	0.42
FeO	0.24	0.23	0.071	0.079	0.078
Optical Characteristics:					
YA (%)	70.0	70.1	72.2	72.0	71.8
TG (nm)	40.6	42.3	63.1	61.6	61.8
Dw (%)	527	525	577	577	576
Pe (%)	2.9	2.5	7.7	7.7	7.6
T _{UV} (%)	6.7	7.2	8.4	7.8	6.9
T ₃₇₀ (%)	19.1	21.1	23.9	22.0	19.5
T ₁₀₅₀ (%)	12.8	14.7	52.4	48.6	49.5
T ₁₂₀₀ (%)	13.1	14.8	51.4	46.9	47.3
λ _p (nm)	1110	1110	1200	1210	1220
Q (J/m ²)	1.5x10 ⁸	1.5x10 ⁸	2.9x10 ⁶	3.6x10 ⁷	2.0x10 ⁸

TABLE 2

	Compara. Example 1	Compara. Example 2	Compara. Example 3	Compara. Example 4
Composition (wt%):				
SiO ₂	71.3	70.5	70.5	70.5
B ₂ O ₃	0	1.0	0	0
Al ₂ O ₃	1.6	1.5	1.5	1.5
MgO	3.64	4.0	4.0	3.46
CaO	7.8	8.0	8.0	8.0
Na ₂ O	12.5	13.0	13.0	13.0
K ₂ O	0.6	0.7	0.7	1.3
CeO ₂	1.6	1.55	1.75	1.7
TiO ₂	0.33	0.03	0.23	0.10
CoO	0	0	0.0011	0.0017
SeO	0	0	0.0011	0.0011
T-Fe ₂ O ₃	0.63	0.7	0.35	0.42
FeO	0.17	0.18	0.054	0.045
Optical Characteristics:				
YA (%)	73.0	71.8	72.8	73.0

TABLE 2 (continued)

	Compara. Example 1	Compara. Example 2	Compara. Example 3	Compara. Example 4
Optical Characteristics:				
TG (nm)	45.2	44.7	66.9	68.9
Dw (%)	534	524	578	578
Pe (%)	3.1	2.5	7.6	8.2
T _{UV} (%)	7.7	8.1	9.2	8.6
T ₃₇₀ (%)	22.5	24.0	26.1	25.1
T ₁₀₅₀ (%)	17.9	17.3	60.1	64.8
T ₁₂₀₀ (%)	19.2	18.4	60.7	65.9
λ_p (nm)	1050	1050	1050	1050

[0043] As is apparent from the results in Tables 1 and 2 above, every sample according to the present invention has an absorption peak wavelength (λ_p) of FeO shifted to the long wavelength side as compared with the corresponding comparative sample. As a result, the proportion of the FeO's absorption reaching to the visible light region is decreased, and the decrease in visible light transmission (YA) is smaller than the decrease in total solar energy transmission (TG) due to ultraviolet irradiation. It is thus understood that the glass exhibiting a visible light transmission comparable to and an infrared absorbing power higher than those of the nonirradiated glass.

[0044] The infrared radiation absorbing glass according to the present invention has an infrared absorption peak at a wavelength of 1,100 nm or longer and therefore exhibits excellent infrared absorbing power and a high visible light transmission.

[0045] Further, due to such high infrared absorption power and visible light transmission, the infrared radiation absorbing glass of the present invention is excellent in reducing the load of air conditioning while securing a comfortable visual field when applied to windows of vehicles, e.g., automobiles, or buildings.

[0046] While the invention has been described in detail and with reference to specific examples thereof, it will be apparent to one skilled in the art that various changes and modifications can be made therein without departing from the scope thereof.

Claims

1. A method for shifting the absorption peak wavelength in the wavelength range 900-1600 nm of an infrared absorbing glass from less than 1100 nm to 1100 nm or longer, comprising the step of irradiating the glass which comprises by wt. %:

65 to 80% SiO₂,
 0 to 5% Al₂O₃,
 0 to 10% MgO,
 5 to 15% CaO,
 10 to 18% Na₂O,
 0 to 5% K₂O,
 5 to 15% MgO + CaO,
 10 to 20% Na₂O + K₂O,
 0 to 5% B₂O₃,
 0.05 to 1.0% total iron oxide (T-Fe₂O₃) in terms of Fe₂O₃,
 0.05 to 2.0% CeO₂,
 0 to 1.0% TiO₂,
 0 to 0.005% CoO, and
 0 to 0.005% Se,

with ultraviolet light to increase the content of FeO in the irradiated glass by reducing Fe(III) to Fe(II), the UV irradiated glass thereby comprising 0.02 to 0.5 wt. % FeO in terms of Fe₂O₃.

2. A method according to Claim 1, wherein the infrared absorbing glass comprises, in % by weight:

0.2 to 0.6% T-Fe₂O₃ in terms of Fe₂O₃;

0.02 to 0.3% FeO in terms of Fe₂O₃;

0.2 to 2.0% CeO₂; and

0.0005 to 0.005% Se.

wherein the relationship between the transmission at 1,050 nm (T_{1050}) and the transmission at 1,200 nm (T_{1200}) is $T_{1050} > T_{1200}$.

3. A method according to Claim 1, wherein the infrared absorbing glass comprises, in % by weight, 0.05 to 3% total of one or more components selected from CeO₂, Sb₂O₃, As₂O₃, SnO, and Cu₂O.

Patentansprüche

1. Verfahren zur Verschiebung der Wellenlänge des Absorptionspeaks im Wellenlängenbereich 900 bis 1.600 nm eines Infrarot absorbierenden Glases von weniger als 1.100 nm auf 1.100 nm oder länger, umfassend den Schritt der Bestrahlung des Glases, das in Gew.% umfasst:

65 bis 80 % SiO₂,

0 bis 5 % Al₂O₃,

0 bis 10 % MgO,

5 bis 15 % CaO,

10 bis 18 % Na₂O,

0 bis 5 % K₂O,

5 bis 15 % MgO + CaO,

10 bis 20 % Na₂O + K₂O,

0 bis 5 % B₂O₃;

0,05 bis 1,0 % Eisenoxid insgesamt (T-Fe₂O₃), bezogen auf Fe₂O₃,

0,05 bis 2,0 % CeO₂,

0 bis 1,0 % TiO₂,

0 bis 0,005 % CoO und

0 bis 0,005 % Se,

mit Ultraviolettlicht zur Erhöhung des Gehalts von FeO im bestrahlten Glas durch Reduktion von Fe(III) zu Fe(II), wobei das UV-bestrahlte Glas hierdurch 0,02 bis 0,5 Gew.% FeO, bezogen auf Fe₂O₃, umfasst.

2. Verfahren gemäß Anspruch 1, wobei das Infrarot absorbierende Glas in Gew.% umfasst:

0,2 bis 0,6 % T-Fe₂O₃, bezogen auf Fe₂O₃;

0,02 bis 0,3 % FeO, bezogen auf Fe₂O₃;

0,2 bis 2,0 % CeO₂; und

0,0005 bis 0,005 % Se,

wobei die Beziehung zwischen der Transmission bei 1.050 nm (T_{1050}) und der Transmission bei 1.200 nm (T_{1200}) $T_{1050} > T_{1200}$ ist.

3. Verfahren gemäß Anspruch 1, wobei das Infrarot absorbierende Glas in Gew.% insgesamt 0,05 bis 3 % von einem oder mehreren Komponenten, ausgewählt aus CeO₂, Sb₂O₃, As₂O₃, SnO und Cu₂O umfasst.

Revendications

1. Procédé pour déplacer le pic d'absorption de longueurs d'onde dans la plage de longueurs d'onde 900 - 1600 nm d'un verre absorbant les infrarouges de moins de 1100 nm à 1100 nm ou plus, comprenant l'étape consistant à irradier le verre qui comprend en % en poids :

65 à 80% de SiO_2 ,
 0 à 5% de Al_2O_3 ,
 0 à 10% de MgO ,
 5 à 15% de CaO ,
 10 à 18% de Na_2O ,
 0 à 5% de K_2O ,
 5 à 15% de $\text{MgO} + \text{CaO}$,
 10 à 20% de $\text{Na}_2\text{O} + \text{K}_2\text{O}$,
 0 à 5% de B_2O_3 ;
 0,05 à 1,0% d'oxyde de fer total ($\text{T-Fe}_2\text{O}_3$) en termes de Fe_2O_3 ,
 0,05 à 2,0% de CeO_2 ,
 0 à 1,0% de TiO_2 ,
 0 à 0,005% de CoO , et
 0 à 0,005% de Se ,

par la lumière ultraviolette pour augmenter la quantité de FeO dans le verre irradié en réduisant Fe (III) en Fe (II) ,
 le verre irradié par UV comprenant ainsi 0,02 à 0,5% en poids de FeO en termes de Fe_2O_3 .

2. Procédé selon la revendication 1, dans lequel le verre absorbant les infrarouges comprend, en % en poids :

0,2 à 0,6% de $\text{T-Fe}_2\text{O}_3$ en termes de Fe_2O_3 ;
 0,02 à 0,3% de FeO en termes de Fe_2O_3 ;
 0,2 à 2,0% de CeO_2 ; et
 0,0005 à 0,005% de Se ,

dans lequel la relation entre la transmission à 1050 nm (T_{1050}) et la transmission à 1200 nm (T_{1200}) est $\text{T}_{1050} > \text{T}_{1200}$.

3. Procédé selon la revendication 1, dans lequel le verre absorbant les infrarouges comprend, en % en poids, 0,05 à 3% au total de un ou plusieurs composants choisis parmi CeO_2 , Sb_2O_3 , As_2O_3 , SnO et Cu_2O .